## AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions and listings of claims in the application:

(Currently amended) A method of making a foam comprising:
providing two syringes connected by a connector, wherein syringe one is
charged with a liquid phase and syringe two is charged with a gas phase, syringe one is
charged with the liquid phase and the gas phase, or both syringes are charged with the
liquid phase and the gas phase; and

transferring the liquid phase and the gas phase repeatedly between the syringes to form a foam, wherein

the liquid phase comprises at least one sclerosing agent and

the gas phase consists essentially of gaseous nitrogen present in an amount ranging from 0.0001% to 0.8% by volume and at least one a physiologically acceptable gas mixture comprising 10% to 90% vol/vol carbon dioxide with the remaining gas oxygen.

- 2. (Previously presented) The method of claim 1, wherein the liquid phase and gas phase passing between the syringes is caused to pass through a mesh comprising apertures with a maximum dimension ranging from 1 to 200 micron.
- (Previously presented) The method of claim 2, wherein the maximum dimension ranges from 2 to 50 micron.

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 (Previously presented) The method of claim 2, wherein the maximum dimension ranges from 3 to 20 micron.

- (Previously presented) The method of claim 1, wherein the gas phase is at least 70% by volume oxygen.
- (Previously presented) The method of claim 1, wherein the gas phase is at least 90% by volume oxygen.
- (Canceled)
- 8. (Canceled)
- 9. (Currently amended) A method of making a foam comprising:
- (a) providing a syringe comprising a barrel, a first plunger and a second plunger, the second plunger having an apertured plunger head which is adapted to be movable within the barrel independently of the first plunger, the syringe being charged with a liquid phase and a gas phase; and
  - (b) oscillating the second plunger to form a foam;

wherein

the liquid phase comprises at least one sclerosing agent and

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the gas phase consists essentially of gaseous nitrogen present in an amount ranging from 0.0001% 0.01% to 0.8% by volume and at least one a physiologically acceptable gas mixture comprising 10% to 90% vol/vol carbon dioxide with the

remaining gas oxygen.

10. (Previously presented) The method of claim 9, wherein the apertures in the

second plunger have a maximum dimension ranging from 1 to 200 micron.

11. (Previously presented) The method of claim 9, wherein the apertures in the

second plunger have a maximum dimension ranging from 2 to 50 micron.

12. (Previously presented) The method of claim 9, wherein the apertures in the

second plunger have a maximum dimension ranging from 3 to 20 micron.

13. (Previously presented) The method of claim 9, wherein the gas phase is at

least 70% by volume oxygen.

(Previously presented) The method of claim 9, wherein the gas phase is at

least 90% by volume oxygen.

15. (Canceled)

16. (Canceled)

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17. (Currently amended) A sterile pack comprising:

(a) a syringe charged with at least one liquid sclerosing agent and a gas mixture

consisting essentially of gaseous nitrogen present in an amount ranging from 0.0001%

0.01% to 0.8% by volume and at least one other a physiologically acceptable gas

mixture comprising 10% to 90% vol/vol carbon dioxide with the remaining gas oxygen;

(b) a gas atmosphere inside the pack having substantially the same composition

as the said gas mixture in the syringe.

18. (Canceled)

19. (Canceled)

20. (Previously presented) The sterile pack of claim 17, wherein the gaseous

nitrogen is present in an amount ranging from 0.01% to 0.7% by volume.

21. (Previously presented) The sterile pack of claim 17, wherein the gaseous

nitrogen is present in an amount ranging from 0.01% to 0.6% by volume.

22. (Canceled)

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